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PROTOCOL PACKAGE ACKNOWLEDGEMENT RECEIPT (Form 2.3)

IRB Protocol No. _____ Date: _____

Protocol Title: _____ Sponsor: _____

Type of Submission:
( ) Initial Submission for review
( ) Resubmission for review
( ) Protocol Amendments
( ) Informed Consent Amendments
( ) Other document for review:
Please specify: _____

Principal Investigator: _____ Contact no./ Email: _____

Documents submitted:

Protocol package for clinical trial and/or sponsor-initiated studies:

- ☐ Letter of Application & Full protocol
- ☐ Protocol Summary
- ☐ Investigator's Brochure
- ☐ Data collection form/s
- ☐ Informed Consent form (English, Tagalog, and local dialect (Hiligaynon))
- ☐ Budget
- ☐ Curriculum Vitae of the Principal Investigator and his/her co-investigators
- ☐ GCP Certificate of the Principal Investigator and his/her co-investigators
- ☐ Declaration of No Conflict of Interest of Investigators/Researchers
- ☐ Photocopy of valid PRC Licence
- ☐ GANTT Chart (as necessary)
- ☐ Advertisement, Diary card and other related documents
- ☐ Case report form/s
- ☐ Certificate of Technical Review (as necessary)

Protocol package for researcher-initiated studies:

- ☐ Letter of Application & Full protocol (Chapter 1-3)
- ☐ Executive summary that follows research project proposal format
- ☐ Data collection form/s
- ☐ Informed Consent form (English/Tagalog, and/or local dialect (Hiligaynon))
- ☐ Budget (as necessary)
- ☐ Curriculum Vitae of the PI and co-investigators
- ☐ Declaration of No Conflict of Interest of Investigators/Researchers
- ☐ GCP Certificate (for intervention studies only)
- ☐ Photocopy of valid PRC Licence
- ☐ GANTT Chart (as necessary)
- ☐ Certificate of Technical Review

Type of Research

- ☐ Clinical Trial ___ Phase 1,2,3,4
- ☐ Biomedical Studies
- ☐ Health Operations research
- ☐ Social research
- ☐ Public Health research
- ☐ Others:

Please specify:
_____
_____
_____

Submitted By:

_____
Signature Over Name:
Date:

Received By:

_____
Signature Over Name:
Date: